

FILED

IN THE UNITED STATE DISTRICT COURT  
FOR THE NORTHERN DISTRICT OF ALABAMA  
MIDDLE DIVISION

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U.S. DISTRICT COURT  
N.D. OF ALABAMA

MARGIE REAVES FOWLER, et al,

Plaintiffs,

v.

Civil Action No.: CV-04-PT-712-M

PHARMACIA and UPJOHN COMPANY,  
MCKESSON CORPORATION,  
CHARLIE WATSON, et al,ENTERED  
JUN 24 2004

## MEMORANDUM OPINION

This cause comes on to be heard upon defendant Charlie Watson's ("Watson") motion to dismiss, filed on April 7, 2004, and plaintiffs Margie Reaves Fowler's ("Mrs. Fowler") and Mark Fowler's ("Mr. Fowler") motion to remand, filed on May 7, 2004.

FACTS<sup>1</sup> AND PROCEDURAL HISTORY

Plaintiff Margie Reaves Fowler ("Fowler") is an adult resident of St. Clair County, Alabama. Plaintiff Mark Fowler is her husband. Defendants Pharmacia & Upjohn Company ("P&U Co.") and McKesson Corporation ("McKesson") are corporations doing business in Alabama. Defendant Watson, an employee of McKesson,<sup>2</sup> allegedly "sold and/or distributed" the drug at issue.

<sup>1</sup> The "facts" are as alleged in the complaint.

<sup>2</sup> The complaint, this court notes, does not allege McKesson Corporation's specific role regarding Depo Provera (also known as medroxyprogesterone). However, defendant Watson's affidavit attached to the notice of removal stated: "My employer, McKesson Medical-Surgical, did not manufacture Depo-Provera Contraceptive Injection. McKesson Medical Surgical was only a distributor of Depo-Provera . . ."



Mrs. Fowler was prescribed and received injections of Depo Provera<sup>3</sup> from June 1999 through December 2001. The injections were given by a licensed health care provider in a clinical setting. Depo Provera was allegedly "manufactured, created, designed, tested, labeled, sterilized, packaged, supplied, marketed, sold, advertised, prescribed, administered and otherwise distributed by the Defendants herein." See Compl. ¶ 7. On February 28, 2002, Mrs. Fowler suffered a stroke, which according to plaintiffs, was proximately caused by Depo Provera. *Id.* at ¶ 9.

On February 27, 2004, plaintiffs filed a complaint in the Circuit Court of St. Clair County, Alabama. The complaint contained the following counts against all defendants: Count One (AEMLD); Count Two (Negligence);<sup>4</sup> Count III (Breach of Express Warranty); Count IV (Breach

<sup>3</sup> Depo Provera is a medication commonly prescribed to women as a contraceptive alternative to "the pill." Depo Provera, taken as an injection, contains a synthetic hormone similar to the natural hormone progesterone and is offered to protect women from pregnancy for three months per injection. See Compl. ¶ 6.

<sup>4</sup> Count Two claims that defendants failed to exercise due care by committing the following acts and omissions:

- a. Failed to adequately and properly test and inspect Depo Provera so as to ascertain whether or not it was safe and proper for the purpose for which it was designed, manufactured or sold
- b. Failed to utilize and/or implement a reasonably safe design in the manufacture of Depo Provera.
- c. Failed to manufacture Depo Provera in a reasonably safe condition for which it was intended;
- d. Failed to adequately and properly warn the Plaintiff purchasing Depo Provera of the risks of complications when used in a manner for which it was intended;
- e. Failed to adequately and properly warn the Plaintiff purchasing Depo Provera of the risks of diseases when used in a manner for which it was intended;
- f. Failed to adequately and properly labeled (*sic*) Depo Provera so as to warn the Plaintiff of the risks of complications;
- g. Failed to adequately and properly label Depo Provera so as to warn the Plaintiff of the risks of complications;

of Implied Warranty); Count V (Damages); Count VI (Unjust Enrichment)<sup>5</sup>; and Count VII (Loss of Consortium by Mr. Fowler). On April 7, 2004, defendants filed a notice of removal, alleging the existence of complete diversity of citizenship between the parties and fraudulent joinder of the non-diverse defendant, Watson. On April 7, 2004, defendant Watson filed the motion to dismiss at issue here.<sup>6</sup> Plaintiffs responded with a motion to remand. The court considers both motions here.

#### RULE 12(b)(6) STANDARD

Rule 12(b)(6) tests the legal sufficiency of a complaint. When considering a Rule 12(b)(6) motion, the court assumes that all factual allegations pled in the complaint are true. *United States*

- h. Manufactured which (sic) constituted a hazard to health;
- i. Manufactured Depo Provera which caused adverse side effects; and
- j. Were otherwise careless and negligent.

See Compl. ¶ 20.

<sup>5</sup> Specifically, Count VI alleges that defendants have profited and benefitted from plaintiff's use of Depo Provera. Additionally, Count VI alleges:

Defendants . . . have voluntarily accepted and retained these profits and benefits, derived from the Plaintiff, with full knowledge and awareness that, as a result of Defendants' . . . fraud and other conscious and intentional wrongdoing, Plaintiff did not receive a product of the quality, nature or fitness that had been represented by Defendants . . . or that Plaintiff, as a reasonable consumer, expected.

By virtue of the conscious wrongdoing alleged in this Complaint, Defendants . . . have been unjustly enriched at the expense of the Plaintiffs, who are entitled to in equity, and hereby seek the disgorgement and restitution of Defendants' . . . wrongful profits, revenue, and benefits, to the extent, and in the amount, deemed appropriate by the Court . . .

*Id.* ¶¶ 35-36.

<sup>6</sup> The court notes that Watson is the only defendant filing the motion to dismiss at issue here. Defendant P&U Co. submitted the response in opposition to plaintiffs' motion to remand. See *infra*.

*v. Gaubert*, 499 U.S. 315, 327, 111 S. Ct. 1267, 113 L. Ed. 2d 335 (1991). All factual allegations are to be construed in the light most favorable to the plaintiff. *Brower v. County of Inyo*, 489 U.S. 593, 598, 109 S. Ct. 1378, 103 L. Ed. 2d 628 (1989). Dismissal under Rule 12(b)(6) is appropriate “only if it is clear that no relief could be granted under any set of facts that could be proved consistent with the allegations” of the complaint.” *Rendon v. Valleycrest Prods., Ltd.*, 294 F.3d 1279, 1282 (11th Cir. 2002) (citing *Hishon v. King & Spalding*, 467 U.S. 69, 73 (1984)).

#### MOTION TO REMAND STANDARD

Federal courts are courts of limited jurisdiction. See *Russell Corp. v. American Home Assurance Co.*, 264 F.3d 1040, 1050 (11th Cir. 2001). Therefore, federal courts have power to hear only those cases that they have been authorized to hear by the Constitution or by Congress. See *Kokkonen v. Guardian Life Ins. Co. of America*, 511 U.S. 375, 377 (1994). The limited nature of federal court jurisdiction has caused the Eleventh Circuit to favor remand of removed cases where federal jurisdiction is not absolutely clear. *Russell Corp.*, 264 F.3d at 1050. The removal statute is to be construed narrowly with doubt construed against removal. See *Shanrock Oil & Gas Corp. v. Sheets*, 313 U.S. 100, 107-09 (1941).

A case may be removed to federal court only if the case could have been brought originally in federal court pursuant to the court’s diversity or federal question jurisdiction. See 28 U.S.C. § 1441(a). However, diversity will not support removal jurisdiction if any properly joined defendants are citizens of the state in which the suit was originally filed. See 28 U.S.C. § 1441(b). The determination of whether federal jurisdiction exists must be made on the face of the plaintiff’s well-pleaded complaint. *Pacheco De Perez v. AT & T Co.*, 139 F.3d 1368, 1373 (11th Cir. 1998). An anticipated or even inevitable federal defense generally will not support removal. *Id.* at 1373 (citing

*Caterpillar, Inc. v. Williams*, 482 U.S. 386, 392-93 (1987)). The burden of establishing federal jurisdiction is placed on the defendant, with all doubts resolved in favor of remand, *Diaz v. Sheppard*, 85 F.3d 1502, 1505 (11th Cir. 1996). When multiple defendants are involved, all defendants must consent to removal. *Russell Corp.*, 264 F.3d at 1050.

### ARGUMENTS

#### MOTION TO DISMISS

##### I. Defendant Watson's Motion

Watson submits that no cause has been stated against him under Alabama law. Relying on *In re Rezulin Products Liability Litigation*, 133 F. Supp. 2d 272, 287-88 (S.D.N.Y. 2001) (predicting Alabama law), Watson argues, a plaintiff patient cannot state a cause of action against a sales representative (or account manager like defendant) of a distributor of a prescription drug. Watson further relies on the Alabama Supreme Court's decision in *Walls v. Alpharma USP, Inc.*, 2004 WL 406759 (Ala. March 5, 2004).

In support of dismissal, Watson relies on his own affidavit, which avers that he was not a manufacturer of Depo-Provera and that he was never involved in the manufacture, development, or testing of the drug. Watson alleges that he has not had dealings with either of the plaintiffs.

Watson's affidavit further provides: Watson has "not made any statements to the general public or participated in any advertising or promotion to the general public concerning Depo Provera . . ."; Watson was not a physician or pharmacist and thus never prescribed or filled a prescription for Depo Provera; As an employee of a distributor, Watson's role was taking orders from physicians' offices; "If their order included a request for Depo-Provera . . . this product, with the information as packaged by the manufacturer, was shipped with the order to their [the physicians'] offices"; and

Watson was not a "seller" of Depo Provera

**A. Counts One and Two - AEMLD and Negligence**

A threshold element of recovery for an AEMLD claim, Watson contends, is showing that defendants "manufactured and/or sold the allegedly defective product." See *Turner v. Azulea Box*, 508 So. 2d 535, 254 (Ala. 1987); *Atkins v. Am. Motors Corp.*, 335 So. 2d 134 (Ala. 1976). Courts from other jurisdictions interpreting Alabama product liability tort theories, Watson claims, have held that no cause of action is stated against sales representatives since they are not "sellers." See *In Re Rezulin*, supra, at 287-88 (S.D.N.Y. 2001). See also *Andrews, et al. v. Bayer Corp., et al; In re Baycol Products Litigation*, MDL No. 1431, slip op. 4 (D. Minn. March 26, 2004) (attached as Exhibit B).

These courts considered the purpose of the AEMLD in analyzing the potential liability of sales representatives of drug manufacturers. *In Re Rezulin* stated: "The AEMLD is founded on 'broader moral notions of consumer protection and on economic and social grounds, placing the burden to compensate for loss incurred by defective products on the one best able to prevent the distribution of these products.'" 133 F. Supp. 2d at 287 (quoting *Atkins* at 139). Furthermore, the *Rezulin* court found: "The sales representative joined in the Alabama case neither manufactured, sold nor supplied Rezulin (the prescription drug at issue in the case). Rather, he was an 'agent of the manufacturer and seller.'" *Id.* at 287-288. Here, Watson repeats, he was neither the manufacturer nor seller of Rezulin. Furthermore, Watson asserts, he is even further removed from liability than the sales representative in *Rezulin* since he was only the distributor's agent.<sup>7</sup>

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<sup>7</sup> Watson again quotes *Rezulin*: "As a corporate employee, he [the representative] was 'not the one best able' to prevent the sale of defective drugs. In light of the Alabama Supreme Court's clear explanation of the AEMLD's scope and purpose, there is no reasonable basis for

Regarding the negligence claims, Watson asserts, they should be dismissed for the same reason, i.e., he was neither the "manufacturer" nor "seller" of Depo Provera. See *Norton Co. v. Harrelson*, 176 So. 2d 18, 20 (Ala. 1965).<sup>8</sup> Additionally, Watson argues, since the negligence count contains the language of the AEMLD, i.e., "It could have been reasonably anticipated by the Defendants . . . that said product would become inherently or imminently dangerous to human life or health when put to its intended, ordinary and customary use," it is redundant with the AEMLD count. Alabama courts have found that when two counts are redundant, the negligence claim is not considered to constitute a separate cause of action. See *Veal v. Teleflex, Inc.*, 586 So. 2d 188, 191 (Ala. 1991).<sup>9</sup>

Watson relies on the *Walls* decision as supporting his position. See Exhibit C. In *Walls*, the plaintiff sued her pharmacist for failure to warn of foreseeable injuries from the use of the prescription drug he dispensed to her. The Northern District of Alabama certified the following question to the Alabama Supreme Court: "Does a pharmacist have a duty to warn of foreseeable supposing that it would impose liability on the sales representatives in this case."

<sup>8</sup> The *Norton* court stated:

This doctrine [of manufacturer's liability] is applicable in a limited number of situations. The defendant must be either the manufacturer or seller of the injury-producing article. There is no privity of contract between the defendant and the injured plaintiff. At the time complained of the article must have been applied to the use for which it was manufactured and sold and that use must be in the usual and customary manner. Where these circumstances exist the manufacturer or seller will be liable for an injury proximately resulting from the use of the article but only where the article is inherently or imminently dangerous to human life or health, or becomes so when put to its intended use in the proper manner. This liability arises from either the negligent manufacture of the article or negligence in selling it.

<sup>9</sup> This court notes that *Veal* does so suggest.



injuries from the use of a prescription drug he/she is dispensing under AEMLD, common-law negligence or other Alabama law?"

In *Walls*, the pharmacist had direct contact with the plaintiff and had directly sold the prescription drug to the plaintiff. Even in that situation, Watson contends, the court applied the learned intermediary doctrine and held that the pharmacist had no duty to warn a customer or any other ultimate customer of the risk or potential side effects of the prescription drug. The Supreme Court observed that

where prescription drugs are concerned the manufacturer's duty to warn is limited to an obligation to advise the prescribing physician of any potential dangers that may result from the drug's use. This special standard for prescription drugs is an understandable exception to the Restatement's general rule that one who markets goods must warn foreseeable ultimate users of dangers inherent in his products. See Restatement (Second) of Torts, Section 388 (1965).

*Walls* at \*3 (citing *Reyes v. Wyeth Laboratories*, 498 F.2d 1264 at 1276)(citations omitted). The *Walls* court further noted language from other cases that to impose a duty to warn on a pharmacist would "intrude on the doctor-patient relationship and would force the pharmacist to practice medicine without a license." *Id.* at \*4.<sup>10</sup>

Defendant concludes: "The rationale which the Alabama Supreme Court followed in holding that prescription drugs are an exception to the Restatement's general rule certainly is even more applicable in the case at bar. If the manufacturer's duty to warn flows only to the physician and if other parties would be liable for interfering in the physician-patient relationship should advice be given to the patient, then it is abundantly clear that Charles Watson cannot be subject to potential liability under Alabama law."

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<sup>10</sup> This court does not find *Walls* to be significantly apt here.



**B. Counts Three and Four - Breach of Express and Implied Warranties<sup>11</sup>**

According to Watson, a breach of warranty claim (whether express or implied) arises exclusively against a product's "seller." See Ala. Code §§ 7-2-313(1), 7-2-314(1), and 7-2-315. The Alabama courts, Watson argues, have affirmed this principle. See, e.g., *Rutledge v. Arrow Aluminum Indust.*, 733 So. 2d 412, 417 (Ala. Civ. App. 1998). The *Rutledge* court found:

With regard to Rutledge's AEMLD and breach of implied warranty of fitness claims against Foshee Builders, it is undisputed that Foshee Builders bought the sliding glass door and that a subcontractor installed the door. Rutledge failed to present any evidence that Foshee Builders is in the business of selling sliding glass doors. Therefore, we conclude that Foshee is not a seller within the meaning of the AEMLD or § 7-2-103 and that the trial court properly entered a summary judgment in favor of Foshee Builders on Rutledge's AEMLD and breach of implied warranty of fitness claims.

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<sup>11</sup> To the extent the warranty claims are redundant with AEMLD claims and based on the allegation that the drug was unreasonably dangerous, defendant argues, these claims are due to be dismissed due to the distinction between tort and UCC causes of action. According to defendant, whether Depo-Provera is unreasonably dangerous is not properly addressed in a warranty claim, only in an AEMLD claim. See *Yarbrough v. Sears, Roebuck & Co.*, 628 So. 2d 478 (Ala. 1993), which found:

The Yarbroughs' claim of a breach of the implied warranty of merchantability is to the effect that the kerosene heater was unreasonably dangerous and therefore could not be merchantable. "Such an argument ignores the clear distinction between causes of action arising under tort law and those arising under the U.C.C. as adopted in Alabama." *Shell v. Union Oil Co.*, 489 So.2d 569, 571 (Ala. 1986). Whether the kerosene heater was unreasonably dangerous is not a question properly addressed in a claim alleging breach of warranty under the U.C.C., but it could be, and was, properly raised in a claim under the AEMLD.

Compare *Spain v. Brown & Williamson Tobacco Corp.*, 2003 WL 21489727 (Ala. 2003) (distinguishing *Yarbrough*). In addressing the certified question from the Eleventh Circuit about the implied warranty of merchantability, the court distinguished *Yarbrough* based on the failure to allege that the product was not fit for the ordinary purpose. The *Spain* court then held: "[A] claim alleging breach of an implied warranty of merchantability is separate and distinct from an AEMLD claim and is viable to redress an injury caused by an unreasonably dangerous product."

Watson reiterates that his position is an account manager of McKesson, a corporation which distributes pharmaceutical products ordered by physicians. As such, Watson asserts, he is not a "seller" for purposes of the U.C.C..

Moreover, Watson argues, an additional reason to dismiss the warranty claim is that Watson had no contact with plaintiffs. Under Alabama law, Watson contends, express warranties arise from affirmative statements of fact. *See* Ala. Code § 7-2-313 (1975). Similarly, Watson argues, an implied warranty cannot arise unless the plaintiff relies on the seller's skill or judgment during the purchase. *See Ex Parte General Motors Corp.*, 169 So. 2d 903, 911 (1999). Since Watson had no contact with plaintiffs, he could not have made affirmative statements or express warranties to them. Additionally, plaintiffs could not have relied on his skill or judgment during their purchase. As a final reason to dismiss plaintiffs' implied warranty claim, Watson asserts, he is not a "merchant with respect to goods of that kind" as required by § 7-2-314. *See Loeb & Co. v. Schreiner*, 321 So. 2d 199 (Ala. 1975); *Huprich v. Bitto*, 667 So. 2d 685 (Ala. 1995).

#### C. Count Five - Unjust Enrichment

Defendant Watson quotes the Alabama Supreme Court in *Mitchell v. H&R Block, Inc.*, 783 So. 2d 812, 817 (Ala. 2000): "[T]he essence of the theories of unjust enrichment .. is that a Plaintiff can prove facts showing that Defendant holds money, which in equity and good conscience, belongs to the Plaintiff or holds money which was improperly paid to Defendant because of mistake or fraud." *See also Ammons v. Coffee County*, 716 So. 2d 1227 (Ala. Civ. App. 1998)

First, Watson argues, since plaintiffs have not stated a valid, independent claim against him, they cannot prevail on their unjust enrichment claim. Second, Watson contends, plaintiffs have not alleged in the complaint that Watson benefitted personally from the sale of Depo-Provera or

collected money from plaintiffs in exchange for this product. Watson repeats his contention that he is not a "seller" of the product.

## II. Plaintiffs' Response<sup>12</sup>

### A. Standard of Review

This court must first determine if it has jurisdiction over the complaint. *See Cabalceta v. Standard Fruit Co.*, 883 F.2d 1553, 1557 (11<sup>th</sup> Cir. 1989); *Univ. of South Alabama v. Am. Tobacco Co.*, 168 F.3d 405, 410 (11<sup>th</sup> Cir. 1999). Strict construction of the removal statutes, plaintiffs argue, is required. *See Shamrock Oil & Gas Corp. v. Sheets*, 313 U.S. 100, 109 (1941); *Clay v. Brown & Williamson Tobacco Corp.*, 77 F. Supp. 2d 1220 (M.D. Ala. 1999).

Further, plaintiffs argue, this court must construe all disputed questions of fact and controlling substantive law in favor of plaintiffs on removal. *Coker v. Amoco Oil Co.*, 709 F.2d 1433, 1440-41 (11<sup>th</sup> Cir. 1983) ("In determining whether joinder of a resident party has been fraudulent, a district court evaluates the factual allegations in the light most favorable to the plaintiff.").

The removing party, plaintiffs assert, bears a heavy burden in establishing fraudulent joinder. The test for determining whether joinder of a defendant has been fraudulent is as follows:

- (1) [L]ook to see whether there is no possibility that plaintiff can establish any cause of action against the resident defendant; and

<sup>12</sup> Plaintiffs note their contemporaneous filing of a motion to remand for lack of jurisdiction. According to plaintiffs, the arguments in support of remand are identical to their response to dismissal. The motion to dismiss and motion to remand, plaintiffs contend, are "really just both sides of the same coin." By plaintiffs' account, if this court grants the motion to remand, the case will be due to be remanded without this court's consideration of the motion to dismiss. On the other hand, plaintiffs contend, if the court denies remand, the court must necessarily have determined plaintiffs had no possibility of establishing any cause of action against the resident defendant, thus requiring the granting of defendant's motion to dismiss.

(2) [L]ook to see whether plaintiff has fraudulently pled jurisdictional facts in order to bring the resident defendant into state court.

*See Cabalceta* at 1561. (Emphasis added) "When considering a motion for remand, federal courts are not to weigh the merits of a plaintiff's claim beyond determining whether it is an arguable one under state law." *Fowler v. Provident Life & Accident Ins. Co.*, 256 F. Supp. 2d 1243, 1247 (N.D. Ala. 2003)(citation omitted). Further, *Fowler* provided: "The plaintiff need not have a winning case against the allegedly fraudulent defendant; she need only have a possibility of stating a valid cause of action in order for the joinder to be legitimate." *Id.* (citation omitted).

#### B. Argument

##### 1. Plaintiffs Have Sufficient Evidence to Support an AEMLD Claim and a Common Law Negligence Claim Against Defendant, Charlie Watson

Plaintiffs tout *Clay v. Brown & Williamson Tobacco Corp.*, 77 F. Supp. 2d 1220 (M.D. Ala. 1999) as analogous. In *Clay*, plaintiffs argue, the court held that plaintiff had an arguable claim against an account manager for defendant pursuant to the AEMLD because the account manager "had superior knowledge to that of the average consumer." *Id.* at 1224. Additionally, the *Clay* court found, additional discovery provided an arguable showing that the account manager "actively participated in the sale and distribution of Brown & Williamson tobacco products," which further bolstered plaintiff's AEMLD claim. In part, the *Clay* court relied on the following rationale from *Seaborn v. R.J. Reynolds Tobacco Co.*, No. 96-T-1540-N (M.D. Ala. 1996):<sup>13</sup>

[Plaintiff] seeks to hold not only R. J. Reynolds liable under the AEMLD, he seeks to hold some of the company's individual employees--Tate, Huffman, McDermott, Hightower, and Hinson--liable as well. "In Alabama, the general rule is that officers or employees of a corporation are liable for torts in which they have personally participated, irrespective

<sup>13</sup> No copy of *Seaborn* has apparently been provided to this court.

of whether they were acting in a corporate capacity." *Ex parte Charles Bell Pontiac-Buick-Cadillac-GMC, Inc.*, 496 So.2d 774, 775 (Ala.1986) (citing *Candy H. v. Redemption Ranch, Inc.*, 563 F.Supp. 505, 513 (M.D.Ala. 1983)); see also *Chandler v. Hunter*, 340 So.2d 818, 822 (Ala.Civ.App. 1976). Obviously, to the extent R.J. Reynolds allegedly violated the AEMLD, it acted through its employees; the company does not employ ghosts. [Plaintiff] should be allowed to pursue these individual defendants, and, if, after discovery, it should turn out that he has named the wrong persons, he should be allowed to make substitutions

In this case and its companion case, *Jenkins v. R.J. Reynolds Tobacco Co. No. 96-T-1489-N* (M.D. Ala. 1996), plaintiffs note, the court found no evidence of fraudulent joinder. Additionally, plaintiffs rely on the *Clay* court's statement: "Rule 11 recognizes that a Plaintiff may need additional discovery to establish an evidentiary basis for an allegation." *Id.* at 1224 (citing *Sellers v. Foremost Ins. Co.*, 924 F. Supp. 1116 (M.D. Ala. 1996)). The court found that plaintiff had met her burden pursuant to Rule 11 and believed that additional discovery could show that the account manager participated in the tort against the plaintiff due to his position with defendant tobacco company.

Here, plaintiffs assert, Watson is an account manager at McKesson, and Birmingham is part of his sales territory. In this capacity, plaintiffs argue, Watson sold, distributed and supplied medical and surgical equipment, i.e., Depo-Provera, to Mrs. Fowler. See Watson Decl.

Plaintiffs rely on "The Job Description of an Account Manager at McKesson Medical-Surgical" posted on McKesson's website, Pl. Ex. B. The job description, plaintiffs argue, lists the following responsibilities for McKesson Account Manager<sup>14</sup>:

- Selling products or services;
- Performing field promotion work and developing new accounts;
- Demonstrating products and/or services and providing assistance in the

<sup>14</sup> The McKesson website, plaintiffs note, lists openings for account managers in cities including LaCrosse and Madison, Wisconsin; Jacksonville, Florida; Portland, Oregon; etc. Plaintiffs contend that all the job descriptions for the locations listed are identical

- best application of products or services;
- Answering all questions concerning products or services and referring questions as necessary;
- Investigating product/service warranty claims to ensure resolution within marketing policies;
- Contacting prospects and explaining features and merits of products or services offered, utilizing persuasive sales techniques.

*See Pl. Ex. B.*

According to plaintiffs, Watson has superior knowledge of Depo-Provera compared to the average consumer, since his job duties include demonstrating the products/services, providing assistance in the best application of the products/services, and explaining the features and merits of the products/services offered. Moreover, plaintiffs argue, in this case they have stronger support for their claims against Watson than the *Clay* plaintiff since they already have proof of Watson's superior knowledge to that of the average consumer regarding Depo-Provera (as evidenced by the foregoing job description).

The AEMLD claim against Watson, plaintiffs contend, is "particularly strong considering that part of Mr. Watson's job description provides for him to 'demonstrate products and/or services and provide assistance in the best application of product or services.'" Plaintiffs further argue:

In this case, the product Depo-Provera, was used for a purpose or application other than the purpose or application for which it had been approved by the Food and Drug Administration (hereinafter FDA). Therefore, through Mr. Watson's assistance with the best application of Depo-Provera, the plaintiff, Margie Fowler was given Depo-Provera for a purpose other than the purpose for which it had been approved by the FDA.<sup>15</sup>

In the complaint, plaintiffs assert, they clearly alleged that Watson was engaged in the business of marketing, selling, advertising, supplying, and distributing Depo-Provera, that the

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<sup>15</sup> This allegation, the court notes, does not appear to be contained in the complaint.

product was defective and unreasonably dangerous, and that as a result Fowler was injured. Plaintiffs repeat the general content of their AEMLD and negligence claims.<sup>16</sup>

Lastly, plaintiffs argue, a corporation's employees are liable for torts in which they personally participated, even if they were acting in a corporate capacity. See *Ex Part Charles Bell Pontiac-Buick-Cadillac-GMC, Inc.*, 496 So. 2d 774, 775 (Ala. 1986). According to plaintiffs, McKesson could not violate the AEMLD on its own; rather, it had to act through its employees. Moreover, plaintiffs add, Watson committed negligence beyond the bounds of the AEMLD.

## 2. The Defendant's Argument for Removal Lacks Legal Support

Plaintiffs attempt to distinguish *In re Baycol* and *In re Rezulin*. According to plaintiffs, these cases represent the opinions of the Minnesota and New York federal district courts, respectively, interpreting Alabama law and thus should not be treated as controlling or given weight in this case. Further, plaintiffs contend, *In re Baycol* involved affidavits from the non-diverse defendants that they were not sellers, manufacturers, developers, or testers of the drug Baycol, and the reported opinion indicates that the *In re Baycol* plaintiffs had no evidentiary support to contradict these affidavits. On the other hand, plaintiffs argue, they possess evidence to contradict Watson's affidavit. Although in *In re Rezulin*, plaintiffs assert, the sales representatives were held to be fraudulently joined, here Watson is not a sales representative but rather an account manager with superior knowledge. Again, plaintiffs rely on *Clay*, *supra*.

Moreover, plaintiffs contend, defendant mistakenly relies on *Walls v. Alparma USPD, Inc.* According to plaintiffs, *Walls* held that a pharmacist does not have a duty to warn a customer or ultimate customer of risks or side effects pursuant to the learned-intermediary doctrine. However,

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<sup>16</sup> The court has summarized the complaint *supra*.



plaintiffs argue, *Walls* neither limits an account manager's duty to warn nor forecloses ABMLD/common law negligence claims against an account manager with superior knowledge and active involvement in the sale/distribution of a product. Furthermore, plaintiffs contend, the pharmacist in *Walls* did not work directly for the manufacturer, distributor, supplier, advertiser, or seller of the drug made the basis of the lawsuit, unlike Watson, who works directly for McKesson (which allegedly distributed, supplied, advertised and sold Depo-Provera )

### III. Defendant Watson's Reply

First, Watson argues, plaintiffs' reliance on *Clay v. Brown & Williamson Tobacco Corp* is misplaced. While *Clay* involved cigarettes and a defendant who worked for the actual manufacturer, the Fowler case involves a prescription medication and a defendant who worked for a distributor. Further, Watson reminds the court, the Alabama Supreme Court recently applied the learned-intermediary doctrine to pharmacists in *Walls*, *see supra*. According to Watson, *Walls* expanded and reinforced the exception to ABMLD liability in cases arising from the use of prescription medications where the only duty to warn runs from the manufacturer to the patient's doctor.<sup>17</sup> *Walls*, Watson argues, clarified that the learned intermediary doctrine forecloses the existence of a duty to

<sup>17</sup> In *Stone v. Smith, Kline & French Laboratories*, 731 F.2d at 1575 (11<sup>th</sup> Cir. 1984), Watson notes, the Eleventh Circuit certified to the Alabama Supreme Court the question of whether an adequate warning to the prescribing physician but not to the ultimate consumer was sufficient as a matter of law. The Alabama Supreme Court in *Stone v. Smith, Kline & French Laboratories*, 447 So. 2d 1301 (Ala. 1984) adopted the Fifth Circuit's learned-intermediary doctrine, which held that pharmaceutical companies who were selling prescription drugs only had a duty to warn the prescribing doctor. According to Watson, the *Walls* court cited with approval language from the *Stone* opinion that imposing a duty to warn on a pharmacist would intrude on the doctor/patient relationship and force the pharmacist to practice medicine without a license, and such reasoning should apply with equal force to Watson. Furthermore, Watson notes, *Walls* also cited language from other courts to the effect that it would be illogical to impose a greater duty on the pharmacist than on the manufacturer. By defendants' account, this same argument should apply to the account manager of a distributor.

warn, and thereby any ABMLD and negligence claim, against someone other than the manufacturer.

As in the instant case, *Walls* involved injuries resulting from a prescription drug. According to Watson, the fact that the product could only be obtained by prescription from a licensed physician led the Supreme Court to apply the learned-intermediary doctrine. Watson again quotes *Walls* and criticizes plaintiffs' position that *Walls* is inapplicable.<sup>18</sup>

In the instant case, Watson argues, the complaint alleges that Depo-Provera was commonly prescribed and that plaintiff was given injections by a licensed health care provider in a clinical setting. Plaintiffs' representation to the court that Watson sold Depo-Provera "to the plaintiff in this case," Watson contends, is false, since his affidavit confirms that he never had contact or dealings with the Fowlers, that he is not a physician or pharmacist, and that he never prescribed or filled a prescription for Depo-Provera. As an employee of the distributor, Watson alleged, his role was limited to taking orders from physicians' offices then shipping any orders for Depo-Provera. According to Watson, he was not involved in the doctor's decision to administer or prescribe Depo-Provera to Mrs. Fowler or in the sale of the product to the plaintiffs. Moreover, Watson points out, Watson has confirmed that he has "not made any statements to the general public or participated in any advertising or promotion to the general public concerning Depo-Provera."

Watson again asserts that the *Walls* rationale regarding prescription drugs as an exception to the Restatement's general rule "is even more applicable in the case at bar" and reasons as follows: "If the manufacturer's duty to warn flows only to the learned-intermediary physician and if other parties, without the medical education or knowledge of the medical history of the patient, would be liable for interfering in the physician-patient relationship should advice be given to the patient, then

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<sup>18</sup> As indicated, this court feels that defendant over-emphasizes the significance of *Walls*.

it is abundantly clear that .. Watson, as an account manager of a *distributor*, owed no duty to the Plaintiffs and cannot be subject to potential liability under Alabama law.”

This court, Watson argues, should not hold that he owed a duty to consumers he had never met and had no way to meet simply because his employer’s website states that he is to demonstrate products and provide assistance. Any such holding, Watson contends, would be contrary to the AEMLD’s purpose stated in *Atkins v. American Motors Corporation*, 335 So. 2d 134, 139 (Alabama 1976): “[The AEMLD is founded on] broader moral notions of consumer protection and on economic and social grounds, placing the burden to compensate for loss incurred by defective products on the one best able to prevent the distribution of these products.” As neither the manufacturer nor seller of Depo-Provera, Watson argues, he is not the “one best able to prevent the distribution of the product.” See *In Re Rezulin* at 287-88 (S.D.N.Y. 2001).<sup>19</sup>

Despite plaintiffs’ argument to the contrary, Watson asserts, *In re Rezulin* and *In re Baycol*, see *supra*, should be given weight, as both federal courts were applying Alabama law. Watson reminds this court of the *Erie* rule, i.e., federal courts exercising diversity jurisdiction must apply the law of the state as interpreted by the state’s highest court, and in the absence of state court precedent, the federal court must ascertain and apply state law as the court would if faced with a similar case. Unlike *Clay* (plaintiffs’ supporting case), Watson argues, *In re Rezulin* and *In re Baycol* directly address AEMLD and negligence claims against sales representatives in cases involving prescription drugs. Notably, Watson asserts, plaintiffs have offered nothing to contradict the holdings of these cases except Watson’s position as an account manager rather than a sales

<sup>19</sup> A threshold element of recovery under AEMLD, Watson repeats, is that plaintiff must prove that the defendant “manufactured and/or sold the allegedly defective product.” See *Turner v. Azalea Box Co.*, 508 So. 2d 253, 254 (Ala. 1987).